## AMENDMENT TO THE CLAIMS

- (Currently Amended) An EPO production system comprising:

   a DNA encoding EPO;
   a vector comprising an HCMV MIEP promoter for receiving the DNA; and
   an avian cella QT-VC for harboring the vector.
- 2. (Canceled)
- 3. (Canceled)
- 4. (Original) The EPO reproduction system of claim 1, wherein the DNA is a genomic DNA encoding EPO.
- 5. (Previously Amended) The EPO production system of claim 1, wherein the DNA encoding EPO is SH (SEQ ID NO: 5).
- 6. (Canceled)
- 7. (Currently Amended) A method of producing EPO comprising the steps of: inserting a DNA encoding an EPO into a vector comprising an HCMV MIEP promoter; transfecting the vector into an avian a QT-VC cell; and culturing the transfected avian cellQT-VC in media.
- 8. (Canceled)
- 9. (Canceled)
- 10. (Original) The method of claim 7, wherein the DNA encoding EPO is a genomic DNA.
- 11. (Previously Amended) The method of claim 7, wherein the DNA encoding the EPO is SH (SEQ ID NO: 5).

Claims 12-14: (Canceled)

- 15. (Currently Amended) An avian cell as a host for expressing EPO by controlling an HCMV-MIEP promoter A quail fibrosarcoma line QT-VC.
- 16. (Canceled)



17. (Currently Amended) The avian cell of claim 15, wherein the avian cell is QT-VCA recombinant cell generated by transfecting a vector comprising a DNA encoding EPO that is under control of HCMV MIEP into a QT-VC cell.

Claims 18-21: (Canceled)